

**K252553 Facial & Body Beauty Device (INIA-BD001, INIA-BD002, INIA-BD003, INIA-BD004, F2C15, F3222 Pro, F5516, F3606, F2210 Pro, F5808, INIA-ED001, INIA-ED002, INIA-BLD001, E1507)**Oct 9, 2025  
57 days to decisionK252553 · Product code: **NFO** · Neurology  
Source: <https://www.510kdatabase.net/k252553/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Transcutaneous Electrical, Aesthetic Purposes (NFO)
Date received	Aug 13, 2025
Decision date	Oct 9, 2025
Days to decision	57 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Shenzhen Jianchao Intelligent Technology Co., Ltd.</b>
Location	Shenzhen, CN
Contact	Fred Li
510(k) history	4 submissions · 4 cleared · 2023-2026

**REGULATORY CONSULTANT**

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Consulting firm	<b>Feiying Drug &amp; Medical Consulting Technical Service Group</b>
Contact	Riley Chen

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)510k Database - [www.510kdatabase.net](https://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k252553/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](https://accessdata.fda.gov). - Generated May 14, 2026